

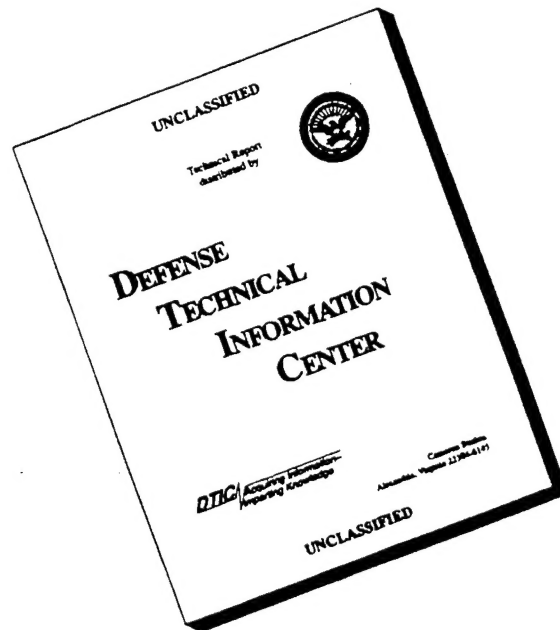
REPORT DOCUMENTATION PAGE

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| 6. AUTHOR(S) Thomas Huntington Brown | | | | |
| 7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Yale University, Grants and Contracts Administration, 12 Prospect Place, New Haven, CT 06511-3516 Attn: Ms Sally Tremaine, Associate Director | | | 8. PERFORMING ORGANIZATION REPORT NUMBER | |
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| 13. ABSTRACT (Maximum 200 words). The near-term goal of the project was to create models of cortical neurons and to implement these on a relatively fast platform. This was done. These models have been used to create insights into the computational differences between neurons and the processing elements typically used in connectionistic studies. The longer-term goal was to abstract these computations into a more efficient form, implement them into learning circuits, and ultimately figure out how to imbed these into low-power, reliable circuit-level VLSI. We have succeeded in the abstractions, and have begun to implement these into circuits that learn and encode time. We are exploring VLSI options. | | | | |
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January 19, 1996

Monique B. Dillon
Procurement Technician
Office of Naval Research
495 Summer Street, Room 103
Boston, MA 02210-2109


RE: N00014-90-J-4136

Dear Ms. Dillon:

Enclosed please find a copy of the Final Technical Report for the referenced grant which was submitted in September 1995 to the program director. Also enclosed is a copy of the Final Invention Statement which was sent to your office in October 1995.

A copy of the technical report is also sent forwarded to the Defense Technical Information.

Cordially,


Sally S. Tremaine
Assistant Director

Enclosures

cc: Defense Technical Information Center (w/enc.)

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DRUG QUALITY IMPROVEMENT

September 7, 1995

Thomas M. McKenna
Office of Naval Research
Code 1142/BI
Division of Cognitive & Neural Sciences
800 North Quincy Street - Room #823
Arlington, VA 22217-5660

Re: MBD
4330
Ser. 479
08 Aug. 1995

Dear Dr. McKenna:

Following is the final technical report on the grant "Self-Organization of Hebbian Synapses on Hippocampal Neurons" (N00014-90-4136). I realize that I normally describe the work in much too great detail, so this time I have pruned it way back.

To reverse-engineer the learning machinery of the brain it is necessary to determine the computations performed by biological neurons and neural circuits and to implement these on a fast platform. This requires defining the device characteristics of the individual neurons and the mechanisms of use-dependent synaptic plasticity.

Our hypothesis has been that learning is a function not only of the training data set but also of the properties of the individual neurons—especially their electrotonic architecture, active membrane properties, and synaptic modification (learning) mechanisms. We have made great strides in unearthing basic neuronal mechanisms and speeding up implementations of neuronal models. This last effort was done in collaboration with computer scientists and a local company that specializes in circuit simulations using fast parallel implementations.

The research has produced numerous publications and abstracts plus presentations at several meetings. There have been major technical, theoretical, and experimental break-throughs. The new technology will result in commercial developments and significant scientific advances. Transitions to industry have been numerous.

The near-term objective of this original proposal was to create models of the key types of hippocampal neurons and be able to implement these on a sufficiently fast platform that we could extract some of their key features. This effort involved a collaboration with Dr. Brenda Claiborne, who was principally responsible for the anatomical aspects of this project.

● As promised, we developed computational models of all of the principal types of hippocampal neurons—CA3 pyramidal neurons, CA1 pyramidal neurons,

and granule cells of the dentate gyrus. Several papers on these model neurons have been published already [1, 2] [3-15] and several more have been submitted or are about to be submitted. A master's thesis was completed on this and is being readied for publication along with three Ph.D dissertations. Together, these will result in 8 - 10 additional publications. In addition, one of my new students is creating models of the inhibitory interneurons, which we know to be critical at the circuit level, which is our current interest (see below).

- Also as promised, we also succeeded in accelerating the implementation of these models, via parallel software that runs on parallel supercomputers or workstation clusters. This is been done using LINDA. In concert with this effort, the Yale Advanced Technology Center, which I founded, hired several relevant scientists to work on aspects of this work. One was Michael Hines, who had made very impressive advances on the development of NEURON, which you probably know now runs on PCs. From my previous progress reports, you realize that we have had significant interactions with and technology transfers to industry.

Much of the initial work centered on the effects of the passive cable properties, but more recent efforts have begun to incorporate various types of nonlinear dynamics and to examine how these interact with various types of synaptic learning rules [16, 17]. There are numerous practical implications for this initial work, which we are continuing under current ONR support. The goal of building silicone brains that learn is no longer far-fetched. Since the ultimate purpose of much work on intelligent systems is to create learning systems, this has been my focus.

- In ONR grant 00014-90-4136, the creation and simulation of model neurons gave me an intimate intuition about how neurons compute. Under the present ONR grant, I have been building on these intuitions to create a completely novel mechanism for associative spatio-temporal learning. Actually, it is not completely novel, as this may be the mechanism that the brain actually uses.

- The transition has been a gradual one—from trying to understand self-organizing systems (the topic of the ONR grant that has just ended) to using this information but also including the objective of accounting for goal-directed or supervised learning. At the synaptic level, of course, we have a good idea that something like a Hebbian mechanism is probably involved [16, 18-27] and this is not hard to implement. The real challenge is to understand how this comes together at the circuit-level. In particular, we need to know how space and time encoded using neuron-like elements.

Until recently, I have been doing these simulations in my head, and they have been working. I now have students collecting the key data and running the circuit simulations to verify my intuitions in this regard, but I am rarely wrong in these matters—that is, usually the simulations simply specify the exact numerical values of the parameters necessary to achieve the intuited result. In the past we have delivered on some key relevant discoveries—demonstrating the existence of Hebbian synapses, showing how they self-organize as a function of their input and electrotonic structure, and devising a whole new theoretical approach to cable theory.

- Now I believe we can bring all of this plus non-linear membrane dynamics into the circuit level in a way that is not hard to implement and can

capture key aspects of supervised spatio-temporal learning in a novel and fascinating manner. I am hoping to have the first simulation of this done within the next year along with the experimental data demonstrating that the postulated mechanisms actually exist in just the right places. Educating myself further in animal learning theory and interactions with the laboratories of Nelson Donnegon and Alan Wagner have been very useful in imposing a top down structure on this collection of devices and microcircuits that we have come to understand.

● Regarding other matters, a final financial report was previously sent under separate cover by our Office of Grants and Contracts. Similarly, they have or will soon send the signed Report of Inventions and Subcontracts form (#882) and the Augmentation Awards form (#A2-2). In regard to the latter, the supported graduate student, Anders Greenwood, has completed his Ph. D work and will do his final dissertation defense on September 26, 1995. He has lined up a Postdoctoral Fellowship at U.C San Francisco Medical School, which will entail working with Robert Malenka and Roger Nicoll, both top neuroscientists.

● Research supported by this grant includes the following as well as several papers that have been submitted or are now being written up for publication. Anticipated publication costs are going to be high, as several of these papers quite long and will include color figures.

1. Brown, T., Zador, A., and Mainen, Z., eds. (1991) *Self-organization of Hebbian synapses in hippocampal neurons*. *Advances in Neural Information Processing Systems*, ed R. Lippmann, et al. Vol. 3. 1991, Morgan Kaufmann. 39-45.

2. Brown, T. H., Zador, A. M., Mainen, Z. F., and Claiborne, B. J. (1992) Hebbian computations in hippocampal dendrites. In *Single neuron computation*, T. McKenna, J. Davis, & S.F. Zornetzer (Eds). Academic Press: San Diego. pp. 81-116.

3. Carnevale, N. T., Tsai, K. Y., Gonzales, R., Claiborne, B. J., and Brown, T. H. (1994) Biophysical accessibility of mossy-fiber synapses in rat hippocampus. *Soc Neurosci Abstr* 20: 715.

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5. Carnevale, N., Tsai, K., Claiborne, B., and Brown, T. (1995) Qualitative electrotonic comparisons of three classes of hippocampal neurons in the rat, In *Neurobiology of Computation*, J. Bower (Ed). Kluwer. pp. 67-72.

6. Kairiss, E. W., Mainen, Z. F., Claiborne, B. J., and Brown, T. H. (1991) Dendritic control of hebbian computations. *Analysis and Modeling of Neural Systems*.

7. Mainen, Z. F., Claiborne, B. J., and Brown, T. H. (1991) A novel role for synaptic computation in the development of cortical lamination. *Soc Neurosci Abstr* 17: 189.

8. O'Boyle, M., Gonzales, R., Brown, T. H., and Claiborne, B. J. (1992) Three-dimensional analyses of granule neurons in the rat dentate gyrus labeled with fluorescent dyes. *Soc Neurosci Abstr* 18: 967.

9. O'Boyle, M. P., Rabimi, O., Brown, T. H., and Claiborne, B. J. (1993) Confocal microscopy allows for improved dendritic diameter measurements and higher input resistances in modeled dentate granules. *Soc Neurosci Abstr* 19: 799.

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13. Zador, A., Koch, C., and Brown, T. H. (1990) Biophysical model of a Hebbian synapse. *Proc Natl Acad Sci (USA)* 87: 6718-6722.
14. Zador, A., Claiborne, B., and Brown, T., eds. (1992) *Nonlinear pattern separation in single hippocampal neurons with active dendritic membrane*. Advances in Neural Information Processing Systems, ed. J. Moody, S. Hanson, & R. Lippman. Vol. 4. 1992, Morgan Kaufmann. 51-58.
15. Zador, A. M., Claiborne, B. J., and Brown, T. H. (1991) Electrotonic transforms of hippocampal neurons. *Soc Neurosci Abstr* 17: 1515.
16. Jaffe, D. B., Fisher, S. A., and Brown, T. H. (1994) Confocal laser scanning microscopy reveals voltage-gated calcium signals within hippocampal dendritic spines. *J Neurobiol* 25: 220-233.
17. Fisher, S. A., Jaffe, D. B., Claiborne, B. J., and Brown, T. H. (1993) Self-organization of Hebbian synapses on a biophysical model of a hippocampal neuron. *Soc Neurosci Abstr* 19: 808.
18. Brown, T. H. and Chattarji, S. (1994) Hebbian synaptic plasticity: Evolution of the contemporary concept. In *Models of Neural Networks II*, J.L. Domany, J.L. van Hemmen, & K. Schulten (Eds). Springer-Verlag: New York. pp. 287-314.
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23. Jaffe, D. B. and Brown, T. H. (1994) Confocal imaging of dendritic Ca^{2+} transients in hippocampal brain slices during simultaneous current- and voltage-clamp recording. *J Microscopy Resch and Tech* 29: 279-289.
24. Jaffe, D. B. and Brown, T. H. (1994) Activity-dependent decay of Ca^{2+} in hippocampal neurons. *Soc Neurosci Abstr* 20: 892.
25. Greenwood, A. C., Xiang, Z., Landaw, E. M., and Brown, T. H. (1993) Likelihood surfaces illuminate uncertainty in quantal analysis. *Soc Neurosci Abstr* 19: 908.
26. Greenwood, A. C., Xiang, Z., Landau, E. M., Kairiss, E. W., and Brown, T., H. (1992) Quantal analysis of plasticity in rat hippocampal mossy-fiber synapses. *Soc Neurosci Abstr* 18: 1350.
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Sincerely,

Thomas H. Brown, Ph. D
Professor of Psychology and Cellular and Molecular Physiology